AMENDMENTS TO THE CLAIMS

Please amend the claims as follows, without prejudice or disclaimer.

1. (Currently amended) A method for inducing an immune response to a tumor antigen in a human being comprising administering a tumor antigen in a first form directly into at least one lymph node and subsequently administering the tumor antigen in a second form ~~that is~~ different from the first form directly into ~~a~~ the at least one lymph node.

2. (Previously Amended) A method according to claim 1 wherein the tumor antigen is selected from the group consisting of CEA, gp100, the MAGE family of proteins, DAGE, GAGE, RAGE, NY-ESO 1, Melan-A/MART 1, TRP-1, TRP-2, tyrosinase, HER-2/neu, MUC-1, p53, KSA, PSA, PSMA, fragments thereof and modified versions thereof.

3. Cancelled

4. (Previously Amended) A method according to claim 1 wherein at least one of said forms is a nucleic acid encoding the tumor antigen and the nucleic acid is selected from the group consisting of viral nucleic acid, bacterial DNA, plasmid DNA, naked DNA, and RNA.

5. (Original) A method according to claim 4 wherein the viral nucleic acid is selected from the group consisting of adenoviral, alphaviral and poxviral nucleic acid.

6. (Original) A method according to claim 5 wherein the poxviral nucleic acid selected from the group consisting of avipox, orthopox and suipox nucleic acid.

7. (Original) A method according to claim 5 wherein the poxviral nucleic acid is selected from the group consisting of vaccinia, fowl pox, canarypox and swinepox nucleic acid.

8. (Original) A method according to claim 5 wherein the poxviral nucleic acid is selected from the group consisting of MVA, NYVAC, TROVAC, and ALVAC nucleic acid.

9. (Previously Amended) A method according to claim 1 wherein at least one of said forms is a nucleic acid encoding the tumor antigen and the nucleic acid is contained in a vector.

10. (Original) A method according to claim 9 wherein the vector is a recombinant virus or bacteria.

11. (Original) A method according to claim 10 wherein the recombinant virus is selected from the group consisting of adenovirus, alphavirus and poxvirus.

12. (Original) A method according to claim 11 wherein the poxvirus is selected from the group consisting of avipox, orthopox and suipox.

13. (Original) A method according to claim 11 wherein the poxvirus is selected from the group consisting of vaccinia, fowlpox, canarypox and swinepox.

14. (Original) A method according to claim 11 wherein the poxvirus is selected from the group consisting of MVA, NYVAC, TROVAC, and ALVAC.

15. (Previously Amended) A method according to claim 1 wherein at least one of said forms is a nucleic acid encoding the tumor antigen and the nucleic acid is contained in a cell.

16. (Previously Amended) A method according to claim 1 wherein at least one of said forms is a nucleic acid encoding the tumor antigen and the nucleic acid is contained in a pharmaceutical composition.

17. (Previously Amended) A method according to claim 1 wherein the tumor antigen is selected from the group consisting of gp100, carcinoembryonic antigen (CEA), a fragment of gp100, a fragment of CEA, a modified version of gp100, and a modified version of CEA.

18. (Previously Amended) A method according to claim 17 wherein the modified version of gp100 comprises at least the sequence IMDQVPFSY (SEQ ID NO: 1) or the sequence YLEPGPVTV (SEQ ID NO:2).

19. (Currently Amended) A method according to claim 17 wherein the modified version of CEA comprises ~~at least the sequence shown in Figure 8 (~~SEQ ID NO:112~~) or the sequence YLSGADLNL (SEQ ID NO:113)~~.

20. Cancelled

21. (Original) A method according to claim 1 wherein the first form is a nucleic acid and the second form is a peptide.

22. (Original) A method according to claim 21 wherein the tumor antigen is selected from the group consisting of CEA, gp100, the MAGE family of proteins, DAGE, GAGE, RAGE, NY-ESO 1, Melan-A/MART 1, TRP-1, TRP-2, tyrosinase, HER-2/neu, MUC-1, p53, KSA, PSA, PSMA, fragments thereof, and modified versions thereof.

23. (Original) A method according to claim 21 wherein the nucleic acid is selected from the group consisting of viral nucleic acid, bacterial DNA, plasmid DNA, naked DNA, and RNA.

24. (Original) A method according to claim 23 wherein the viral nucleic acid is selected from the group consisting of adenoviral, alphaviral and poxviral nucleic acid.

25. (Original) A method according to claim 24 wherein the poxviral nucleic acid selected from the group consisting of avipox, orthopox and suipox nucleic acid.

26. (Original) A method according to claim 25 wherein the poxviral nucleic acid is selected from the group consisting of vaccinia, fowl pox, canarypox and swinepox nucleic acid.

27. (Original) A method according to claim 26 wherein the poxviral nucleic acid is selected from the group consisting of MVA, NYVAC, TROVAC, and ALVAC nucleic acid.

28-31. Canceled

32. (Previously presented) The method of claim 1 wherein both a humoral and cell mediated immune response greater than that produced by subcutaneous immunization are observed.

33. (Previously presented) The method of claim 1 wherein the tumor antigen is not co-administered with an adjuvant.

34-35. Cancelled

36. (New) The method of claim 21 wherein the peptide is selected from the group consisting of SEQ ID NO.: 1, SEQ ID NO.: 2, and SEQ ID NO.: 113.

37. (New) The method of claim 1 wherein the first form of the tumor antigen comprises SEQ ID NO.: 110.

38. (New) The method according to claim 17 wherein the modified version of CEA comprises SEQ ID NO:113.

39. (New) The method of claim 1 wherein the first form of the tumor antigen comprises SEQ ID NO.: 110 and the second form of the tumor antigen is at least one of the peptides SEQ ID NO.: 1 or SEQ ID NO.: 2.